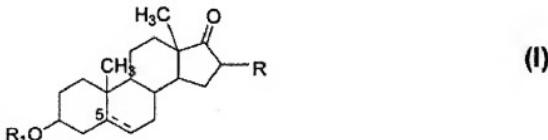


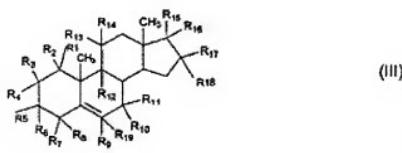
Amendments to the Claims:

1. (Original) A pharmaceutical composition, comprising a pharmaceutically or veterinarianily acceptable carrier, a first active agent and a second active agent effective to treat asthma, chronic obstructive pulmonary disease, or a respiratory or lung disease,

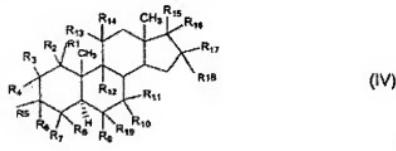
(a) the first active agent is a non-glucocorticoid steroid having the chemical formula



wherein the broken line represents a single or a double bond; R is hydrogen or a halogen; the H at position 5 is present in the alpha or beta configuration or the compound of chemical formula I comprises a racemic mixture of both configurations; and R¹ is hydrogen or a multivalent inorganic or organic dicarboxylic acid covalently bound to the compound; a non-glucocorticoid steroid of the chemical formula



a non-glucocorticoid steroid of the chemical formula



wherein R1, R2, R3, R4, R5, R7, R8, R9, R10, R12, R13, R14 and R19 are independently H, OR, halogen, (C1-C10) alkyl or (C1-C10) alkoxy, R5 and R11 are independently OH, SH, H, halogen, pharmaceutically acceptable ester, pharmaceutically acceptable thioester, pharmaceutically acceptable ether, pharmaceutically acceptable thioether, pharmaceutically

acceptable inorganic esters, pharmaceutically acceptable monosaccharide, disaccharide or oligosaccharide, spirooxirane, spirothirane, -OSO₂R₂₀, -OPOR₂₀R₂₁ or (C₁-C₁₀) alky, R₅ and R₆ taken together are =O, R₁₀ and R₁₁ taken together are =O; R₁₅ is (1) H, halogen, (C₁-C₁₀) alkyl, or (C₁-C₁₀) alkoxy when R₁₆ is -C(O)OR₂₂, (2) H, halogen, OH or (C₁-C₁₀) alkyl when R₁₆ is halogen, OH or (C₁-C₁₀) alkyl, (3) H, halogen, (C₁-C₁₀) alkyl, (C₁-C₁₀) alkenyl, (C₁-C₁₀) alkynyl, formyl, (C₁-C₁₀) alkanoyl or epoxy when R₁₆ is OH, (4) OR, SH, H, halogen, pharmaceutically acceptable ester, pharmaceutically acceptable thioester, pharmaceutically acceptable ether, pharmaceutically acceptable thioether, pharmaceutically acceptable inorganic esters, pharmaceutically acceptable monosaccharide, disaccharide or oligosaccharide, spirooxirane, spirothirane, -OSO₂R₂₀ or -OPOR₂₀R₂₁ when R₁₆ is H, or R₁₅ and R₁₆ taken together are =O; R₁₇ and R₁₈ are independently (1) H, -OH, halogen, (C₁-C₁₀) alkyl or -(C₁-C₁₀) alkoxy when R₆ is H OR, halogen, (C₁-C₁₀) alkyl or -C(O)OR₂₂, (2) H, (C₁-C₁₀) alkyl, ((C₁-C₁₀) alkyl)_n amino-(C₁-C₁₀) alkyl, (C₁-C₁₀) alkoxy, hydroxy-(C₁-C₁₀) alkyl, (C₁-C₁₀) alkoxy-(C₁-C₁₀) alkyl, (halogen)_m (C₁-C₁₀) alkyl, (C₁-C₁₀) alkanoyl, formyl, (C₁-C₁₀) carbalkoxy or (C₁-C₁₀) alkanoyloxy when R₁₅ and R₁₆ taken together are =O, (3) R₁₇ and R₁₈ taken together are =O; (4) R₁₇ or R₁₈ taken together with the carbon to which they are attached form a 3-6 member ring containing 0 or 1 oxygen atom; or (5) R₁₅ and R₁₇ taken together with the carbons to which they are attached form an epoxide ring; R₂₀ and R₂₁ are independently OH, pharmaceutically acceptable ester or pharmaceutically acceptable ether; R₂₂ is H, (halogen)_m (C₁-C₁₀) alkyl or (C₁-C₁₀) alkyl; n is 0, 1 or 2; and m is 1, 2 or 3; or pharmaceutically or veterinarianally acceptable salts thereof; and

(b) the second active agent is a tyrosine kinase inhibitor, delta opioid receptor antagonist, neurokinin receptor antagonis, or VCAM inhibitor.

2. (Original) The pharmaceutical composition of claim 1, wherein the first active agent is a non-glucocorticoid steroid having the chemical formula (I), wherein said multivalent organic dicarboxylic acid is SO₂OM, phosphate or carbonate, wherein M comprises a counterion, wherein said counterion is H, sodium, potassium, magnesium, aluminum, zinc, calcium, lithium,

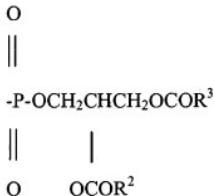
ammonium, amine, arginine, lysine, histidine, triethylamine, ethanolamine, choline, triethanolamine, procaine, benzathine, tromethamine, pyrrolidine, piperazine, diethylamine, sulfatide



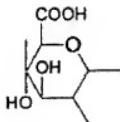
|



or phosphatide



wherein R² and R³, which are the same or different, and are straight or branched (C₁-C₁₄) alkyl or glucuronide



3. (Original) The pharmaceutical composition of claim 2, wherein said first active agent is dehydroepiandrosterone.

4. (Original) The pharmaceutical composition of claim 2, wherein said first active agent is dehydroepiandrosterone-sulfate.

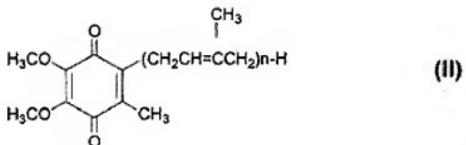
5. (Withdrawn) The pharmaceutical composition of claim 1, wherein said second agent is a tyrosine kinase inhibitor.

6. (Withdrawn) The pharmaceutical composition of claim 1, wherein said second agent is a delta opioid receptor antagonist.

7. (Withdrawn) The pharmaceutical composition of claim 1, wherein the second agent is a neurokinin receptor antagonist.

8. (Withdrawn) The pharmaceutical composition of claim 1, wherein the second agent is a VCAM inhibitor.

9. (Original) The pharmaceutical composition of claim 1, further comprising a ubiquinone or pharmaceutically or veterinarily acceptable salt thereof, wherein the ubiquinone has the chemical formula.



wherein n is 1 to 12

10. (Original) The pharmaceutical composition of claim 1, wherein the pharmaceutical composition comprises particles of inhalable or respirable size.

11. (Original) The pharmaceutical composition of claim 10, wherein the particles are about 0.01 μm in size.

12. (Original) The pharmaceutical composition of claim 10, wherein the particles are about 10 μm to about 100 μm in size.

13. (Original) A kit comprising a delivery device and the pharmaceutical composition of claim 1.

14. (Original) The kit of claim 13, wherein the delivery device is an aerosol generator or spray generator.
15. (Original) The kit of claim 11, wherein the aerosol generator comprises an inhaler.
16. (Original) The kit of claim 15, wherein the inhaler delivers individual pre-metered doses of the formulation.
17. (Original) The kit of claim 15, wherein the inhaler comprises a nebulizer or insufflator.
18. (Withdrawn) A method for reducing the probability of or treating asthma in a subject, comprising administering to a subject in need of such treatment a prophylactically or therapeutically effective amount of the pharmaceutical composition of claim 1.
19. (Withdrawn) A method for reducing the probability of or treating of chronic obstructive pulmonary disease in a subject, comprising administering to a subject in need of such treatment a prophylactically or therapeutically effective amount of the pharmaceutical composition of claim 1.
20. (Withdrawn) A method for treatment of respiratory, lung or malignant disorder or condition, or for reducing levels of, or sensitivity to, adenosine or adenosine receptors in a subject, comprising administering to a subject in need of such treatment a prophylactically or therapeutically effective amount of the pharmaceutical composition of claim 1.
21. (Withdrawn) The method of claim 20, wherein the disorder or condition comprises asthma, chronic obstructive pulmonary disease (COPD), cystic fibrosis (CF), dyspnea, emphysema, wheezing, pulmonary hypertension, pulmonary fibrosis, hyper-responsive airways, increased adenosine or adenosine receptor levels, adenosine hyper-sensitivity, infectious diseases, pulmonary bronchoconstriction, respiratory tract inflammation or allergies, lung surfactant or ubiquinone depletion, chronic bronchitis, bronchoconstriction, difficult breathing, impeded or obstructed lung airways, adenosine test for cardiac function, pulmonary

Appln. No.: 10/698,071

Restriction Requirement Mailed On January 5, 2007

Response To Restriction Filed On March 5, 2007

vasoconstriction, impeded respiration, Acute Respiratory Distress Syndrome (ARDS), administration of adenosine or adenosine level increasing drugs, infantile Respiratory Distress Syndrome (infantile RDS), pain, allergic rhinitis, cancer, or chronic bronchitis.